

## ABSTRACT OF THE DISCLOSURE

Protein kinase C (PKC) has been implicated as a mediator of diabetes-induced vascular proliferation. This study investigated the regulation of PKC $\beta$  gene expression following acute glucose exposure in human vascular smooth muscle cells and in A10 cells, a rat aortic smooth muscle cell line. Western blot analysis showed that PKC $\beta$ II protein levels decreased with high glucose (25mM) while PKC $\beta$ I levels were unaltered. PKC $\beta$  mRNA levels were depleted by 60-75% in hyperglycemic conditions. Quenching of PKC $\beta$  promoter activity by glucose suggested involvement of a carbohydrate response element in the 5' promoter region. Simultaneous cell cycle studies indicated an increase in the percentage of cells going into S phase in high glucose implying that quenching of PKC $\beta$  transcription may be related to cell cycle progression. It was demonstrated that glucose induced post-transcriptional destabilization of PKC $\beta$ II message via a nuclease activity present in the cytosol. The specificity of glucose to post-transcriptionally destabilize PKC $\beta$ II, but not the PKC $\beta$ I, isoform was confirmed in both A10 cells and primary cultures from human aorta.

To further elucidate the intracellular signaling mechanisms, glucose analogs were used to study the pathways by which glucose acted to destabilize PKC $\beta$ II mRNA. Glucose-induced destabilization of PKC $\beta$ II mRNA is independent of the hexosamine or hexokinase pathways. Cycloheximide did not block destabilization of PKC $\beta$ II mRNA by high glucose indicating that the process is independent of translation. Glucose may act

via PKC signaling pathways and may be regulated by serine/threonine phosphorylation/dephosphorylation.

A heterologous chimeric minigene encoding PKC $\beta$ II cDNA subcloned into the p $\beta$ G expression vector comprising the coding sequences of  $\beta$ -globin genomic DNA and 3'UTR and polyadenylation site of bovine growth hormone cDNA was constructed. Half-life analysis indicated a rapid glucose-induced destabilization of  $\beta$ -globin mRNA in p $\beta$ G-PKC $\beta$ II transfected cells. Mobility shift analysis indicated the presence of a glucose-responsive instability element within the exon included in the mature PKC $\beta$ II mRNA in VSMC. UV cross-linking analysis showed a small protein (~10-14 kDa) binding near a stem-loop structure within the PKC $\beta$ II-specific exon. This is a novel finding of a instability element within the PKC $\beta$ II mRNA coding region that is regulated by glucose in aorta smooth muscle cell.